GOOD SENSE PAIN RELIEF- acetaminophen tablet L. Perrigo Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Perrigo Pain Relief Tablets Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- the common cold
- headache
- minor pain of arthritis
- backache
- muscular aches
- toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Do not use

• with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

• if the user has ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if the user

- has liver disease
- is a child with pain of arthritis

Ask a doctor or pharmacist before use if the user is

taking the blood thinning drug warfarin

Stop use and ask a doctor if

- new symptoms occur
- pain gets worse or lasts for more than 10 days (for adults) or 5 days (for children)
- fever gets worse or lasts for more than 3 days
- redness or swelling is present

These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

do not take more than directed (see overdose warning)

adults and children 12 years and over	 take 2 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours do not use for more than 10 days unless directed by a doctor
children 6-11 years	 take 1 tablet every 4 to 6 hours while symptoms last do not take more than 5 tablets in 24 hours do not use for more than 5 days unless directed by a doctor
children under 6 years	ask a doctor

Inactive ingredients

croscarmellose sodium*, povidone, pregelatinized starch, stearic acid *may contain this ingredient

Questions or comments?

1-800-719-9260

Principal Display Panel

Regular Strength
Pain Relief TABLETS
Pain Reliever/Fever Reducer

Acetaminophen

Compare to active ingredient of Tylenol® Tablets 100% SATISFACTION GUARANTEED

100 Tablets - 325 mg Each



Compare to adm Tylenol® Tablets



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Drug Facts (continued)

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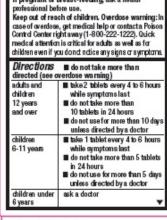
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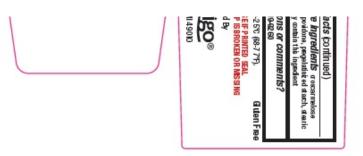
help right away.

Do not use

100 Tablets - 325 mg Each







GOOD SENSE PAIN RELIEF

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0113-0403
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	325 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M28 O L1HH48)		
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
STEARC ACD (UNI. 4EE V/200AF)		

Product Characteristics			
Color	WHITE	Score	no score
Shape	ROUND (beveled edge)	Size	10 mm
Flavor		Imprint Code	325MG;L403
Contains			

Ш	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0113-0403-78	1 in 1 CARTON	08/15/1989	
	1	100 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	08/15/1989	

Labeler - L. Perrigo Company (006013346)

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